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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/940,518	08/29/2001	Loong-Tak Lim	13201.00051	5213
27160 75	90 09/30/2003			
PATENT ADMINSTRATOR KATTEN MUCHIN ZAVIS ROSENMAN 525 WEST MONROE STREET			EXAMINER	
			GAKH, YELENA G	
SUITE 1600 CHICAGO, IL 60661-3693			ART UNIT	PAPER NUMBER
00.100,			1743	ı ſ
			DATE MAILED: 09/30/2003	14

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)				
200	09/940,518	LIM ET AL.				
Office Action Summary	Examiner	Art Unit				
•	Yelena G. Gakh, Ph.D.	1743				
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply						
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely. - If NO period for reply specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). - Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b). Status						
1) Responsive to communication(s) filed on 30 J	<u>uly 2003</u> .					
2a)⊠ This action is FINAL . 2b)□ Thi	s action is non-final.					
3) Since this application is in condition for allowance except for formal matters, prosecution as to the ments is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.						
Disposition of Claims A) Claim(s) 25 27 76 104 and 105 is/are panding in the application						
 4)⊠ Claim(s) 35,37-76,104 and 105 is/are pending in the application. 4a) Of the above claim(s) is/are withdrawn from consideration. 						
5) Claim(s) is/are allowed.						
6)⊠ Claim(s) <u>35,37-76,104 and 105</u> is/are rejected.						
7) Claim(s) is/are objected to.						
8) Claim(s) are subject to restriction and/or election requirement.						
Application Papers						
9) The specification is objected to by the Examiner.						
10) The drawing(s) filed on is/are: a) □ accepted or b) □ objected to by the Examiner.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
11)☐ The proposed drawing correction filed on is: a)☐ approved b)☐ disapproved by the Examiner.						
If approved, corrected drawings are required in reply to this Office action.						
12) The oath or declaration is objected to by the Examiner.						
Priority under 35 U.S.C. §§ 119 and 120						
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).						
a) All b) Some * c) None of:						
1. Certified copies of the priority documents have been received.						
2. Certified copies of the priority documents have been received in Application No						
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 						
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).						
a) ☐ The translation of the foreign language provisional application has been received. 15)☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.						
Attachment(s)						
 Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449) Paper No(s) 7.9 	5) Notice of Informal F	/ (PTO-413) Paper No(s) Patent Application (PTO-152)				

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DETAILED ACTION

1. The Amendment filed on 07/30/03 is acknowledged. Claims 1-34, 36 and 77-100 are cancelled without prejudice. Claims 35, 37-76 and 104-105 are pending in the Application.

Response to Amendment

2. Rejection of claim 76 under 35 U.S.C. 112, second paragraph, is withdrawn in view of the amendment. Rejection of the pending claims over the prior art is modified due to the amendment as following.

Claim Rejections - 35 USC § 103

- 3. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
 - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 4. The factual inquiries set forth in *Graham* v. *John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:
 - 1. Determining the scope and contents of the prior art.
 - 2. Ascertaining the differences between the prior art and the claims at issue.
 - 3. Resolving the level of ordinary skill in the pertinent art.
 - 4. Considering objective evidence present in the application indicating obviousness or nonobviousness.
- 5. This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out

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the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

6. Claims 35, 37-38, 40-42, 44-45, 47-49, 51-54, 63-67 and 104 are rejected under 35 as being unpatentable over Iannacone et al. (US 3,645,696) in view of Moore (US 5,332,548) and Hauser (US Public Health Service Publication, 1965).

Iannacone discloses "a method for the preparation of a chromogenic reagent for aldehydes which is normally unstable upon exposure to air. The invention comprises the dissolving of the chromogenic reagent in a solvent and subsequently pouring the solvent containing the dissolved chromogenic reagent onto a support medium [carrier], while constantly in the presence of an inert atmosphere. The support medium may be dried simultaneously with the adsorption of the chromogenic reagent. Some chromogenic reagents, which may be utilized, are salicylalhydrazone, p-nitrobenzalhydrazone, 2-hydrazinobenzothiazole, 2-hydrazinobenzothiazole-4-nitrobenzenediazonium fluoborate, etc., while 3-methyl-2-benzothiazolinone hydrazone hydrochloride monohydrate (MBTH) is preferred. One of the solvents used to dissolve the chromogenic reagent to further its introduction onto the support medium is deionized and/or distilled water, while any solution, which will dissolve the test reagent, may be utilized. The support medium may consist of any suitable material such as anhydrous aluminum, silica gel, diatomaceous earth, firebrick or polymeric material. Suitable polymeric material includes polyethylene, nylon, etc. and is preferably present in particulate form such as chips or powder. The inert atmosphere may be provided such as by introducing a continuous flow of dry nitrogen gas through the support medium during the adsorption period. The drying process is generally conducted at a temperature less than about 100 °C during the entire preparation of the chromogenic reagent material" (col. 2, lines 5-38). "The MBTH solution in deionized water would generally be a 0.5 to 5 percent solution" (col. 3, lines 43-45). The carrier, which is a support, is inert to the aldehyde-reactive reagent.

Iannacone does not specifically teach applying the reagent-coated carrier to a support.

Moore teaches a method for preparation of "an analytical device for detecting a gaseous or volatile analyte emitted or released from a solid material, and is adaptable to be applied directly to or in contact with a surface of the solid material. The device comprises (a) a substrate

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for adjacent disposition on a surface of the solid material from which the analyte is emitted, said substrate being substantially inert to the analyte, (b) an analyte-reactive component, and (c) an indicator" (col. 4, lines 6-13). "Although the invention is described with particular reference to detecting or determining **formaldehyde** emissions, it should be understood that the invention is also applicable to other gaseous or volatile analytes which meet the criteria of the invention, as for example, isobutane, acetone, butyl acetate, methyl acrylate, isopropyl acetate, methyl ethyl ketone, diisobutyl ketone, **acetaldehyde**, etc." (col. 7, lines 40-47).

"In a preferred embodiment, the substrate comprises a planar strip of high purity paper ... cut from a mother web or roll. Thus, the mother roll can be coated or impregnated first with the analyte-reactive component, as by spraying, dipping, brushing, spreading, printing, etc., and then dried in air or preferably in an inert or non-interfering atmosphere such as nitrogen, and/or dried in a low temperature oven of from about 25° to 90° C, which in any case may be conducted under reduced pressure, although the drying temperature can be higher depending largely on such factors as type of substrate, composition and concentration of the reagent, and drying time. The indicator coating is next applied in a similar manner" (col. 8, lines 28-48). "Suitable substrate materials include natural and synthetic materials, such as cellulose (e.g., wood cellulose, cotton, and rayon), polyolefins (e.g., polyethylene, and polypropylene), nylon, glass fibers, and ceramic-type materials" (col. 8, lines 8-11). "Synthetic materials include synthetic papers, nonwoven polyolefins, glass fibers, glass beads, and such materials as silica gel, activated alumina, and molecular sieves applied as a thin coating on an inert carrier [support] which preferably is transparent such as polyethylene tape" (col. 8, lines 20-25).

"The analyte-reactive component is applied to at least a portion of the substrate typically as a solution as by dipping, spraying, spreading, brushing, or printing" (col. 8, lines 66-68). The adhesive polyethylene tape can be single or double sided depending on the embodiment, as disclosed in column 11.

It would have been obvious for anyone of ordinary skill in the art to slightly modify Iannocone's method of making his indicator by applying the reagent-coated carrier to a support the same way Moore discloses for his indicator for the reasons indicated by Moore, in col. 5, i.e. to provide "interjacent disposition between the substrate [coating] and the solid material" (lines 16-17), as well as to protect the indicator (lines 35-45).

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Iannocone in view of Moore do not specifically disclose a developing solution for use in quantifying an amount of acetaldehyde, in particular ferric chloride.

Hauser discloses MBTH method for determination of aliphatic aldehydes, comprising oxidizing azines resulted from interaction of MBTH with aldehydes by "FeCl₃-sulfamic acid solution to form the blue cationic dye, which can be measured at 628 nm" (Abstract).

It would have been obvious for anyone of ordinary skill in the art to modify Iannocone-Moore's method of making the indicator by providing oxidizing developing solution, ferric chloride in particular, because a step of developing MBTH-aldehyde product with this reagent is a part of the well-known analytical method for quantitative detection of aldehydes, as indicated by Hauser back in 1965.

7. Claim 39 is rejected under 35 U.S.C. 103(a) as being unpatentable over Iannacone in view of Moore and Hauser, as applied to claims 35, 37-38, 40-42, 44-45, 47-49, 51-54, 63-67 and 104 above, and further in view of e.g. Cohen et al. (US 3,649,159).

Iannacone in view of Moore and Hauser do not particularly disclose optimizing pH of solution for promoting solubility of an acetaldehyde-reactive reagent. However it is well-known that solubility of many organic compounds, which are not completely dissolved in water, such as dyes, depends on pH of the solution, as indicated e.g. by Cohen: "in many instances, the optimum pH is dependent upon the solubility of the dye material" (col. 4, lines 30-31).

It would have been obvious for anyone of ordinary skills in the art to optimize pH of the solution to promote solubility of an aldehyde-reactive reagent, disclosed by Iannacone-Moore - Hauser since in many instances it is a dye, which is not completely dissolved in water, as indicated by Cohen.

8. Claims 43, 46 and 50 are rejected under 35 U.S.C. 103(a) as being unpatentable over Iannacone in view of Moore and Hauser, as applied to claims 35, 37-38, 40-42, 44-45, 47-49, 51-54, 63-67 and 104 above, and further in view of the well known prior art, e.g. Meier et al. (EP 885914 A2, Abstract).

Although Iannacone in view of Moore and Hauser do not specifically indicate a membrane, foam and kaolin, it would have been obvious for anyone of ordinary skills to use a membrane, foam or kaolin as suitable carriers, since polymeric membranes and foams are well known conventional carriers along with gels and films, as indicated e.g. in Meier: "gels, films

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and foams may have an extremely high water uptake capacity" (see Advantage), and kaolin (alumina silicate) is a part of ceramic materials indicated by Moore.

9. Claims 55-61 are rejected under 35 U.S.C. 103(a) as being unpatentable over Iannacone in view of Moore and Hauser, as applied to claims 35, 37-38, 40-42, 44-45, 47-49, 51-54, 63-67 and 104 above, and further in view of the well-known prior art, indicated e.g. in Jilla (US 3,932,126).

Although Iannacone in view of Moore and Hauser do not disclose various sources of heat recited in claims 55-61, it would have been obvious for anyone of ordinary skills in the art to use any controllable sources of heat for drying the reagent, such as disclosed e.g. in Jilla: "the initial drying of impregnated polyester material may be carried out in any convenient equipment per se well known in the art, such as a **dielectric heater**, a microwave heater, an infrared heater" (col. 2m lines 35-38), since only controlled temperature is important for the correct conditions of drying so as not to degrade the reagent. A water bath is known to give temperatures less then 100° C, which is in the range disclosed by Iannacone and Moore.

10. Claims 68-74, 76 and 105 are rejected under 35 U.S.C. 103(a) as being unpatentable over Iannacone in view of Moore, Wang (US 4,622,207) and Hauser.

Iannacone teaches a method for making an indicator for testing acetaldehyde by contacting a solution, of 3-methyl-2-benzothiazoline hydrozone (MBTH), particularly 0.5-5% solution in water, with a particulate carrier (e.g. alumina or silica gel, col. 2, line 21) and drying it in a non-reactive atmosphere.

Iannacone does not specifically teach applying a thin layer of the coated particulate carrier to an inert adhesive tape.

Moore teaches a method for preparation of a similar acetaldehyde indicator with a specific structure of attached reagent-coated particular carrier to an inert "polyethylene [adhesive] tape" (col. 8, lines 20-25). The mean emission rate moles detected by the indicator are $0.5 \times 10^{-8} - 1.9 \times 10^{-6}$ CH₂O/cm²/hr, which is equivalent to the detection limit indicated in claim 76.

It would have been obvious for anyone of ordinary skill in the art to slightly modify Iannocone's method of making his indicator by applying the reagent-coated carrier to a support the same way Moore discloses for his indicator for the reasons indicated by Moore, in col. 5, i.e.

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to provide "interjacent disposition between the substrate [coating] and the solid material" (lines 16-17), as well as to protect the indicator (lines 35-45).

Iannacone in view of Moore do not specifically disclose bonding the adhesive tape with a thin layer of the coated particulate carrier to a support.

Wang describes "a typical construction of a reagent device employing filter paper as the carrier matrix material", wherein a carrier matrix with a reagent is affixed by means of "double faced adhesive tape 16 to an insoluble support member, such as organoplastic strip 18" shown in Figure 1 (col. 5, lines 31-35).

It would have been obvious for anyone of ordinary skills in the art to modify Iannacone-Moore's method of making the indicator by adding a step of bonding the adhesive tape with a thin layer of the coated particulate carrier to a support the way it is indicated by Wang, because it provides a simple and conventional structure for the acetaldehyde indicator, as emphasized by Wang.

It would have been obvious for anyone of ordinary skill to use one-sided tape for the same purpose and attach it to the support with additional means, e.g. another adhesive tape, as this is an obvious way for anyone of ordinary skill to attach anything to the surface.

Iannocone in view of Moore and Wang do not specifically disclose a developing solution for use in quantifying an amount of acetaldehyde, in particular ferric chloride.

Hauser discloses MBTH method for determination of aliphatic aldehydes, comprising oxidizing azines resulted from interaction of MBTH with aldehydes by "FeCl₃-sulfamic acid solution to form the blue cationic dye, which can be measured at 628 nm" (Abstract).

It would have been obvious for anyone of ordinary skill in the art to modify Iannocone-Moore-Wang's method of making the indicator by providing oxidizing developing solution, ferric chloride in particular, because a step of developing MBTH-aldehyde product with this reagent is a part of the well-known analytical method for quantitative detection of aldehydes, as indicated by Hauser back in 1965.

11. Claim 75 is rejected under 35 U.S.C. 103(a) as being unpatentable over Iannacone in view of Moore, Wang (US 4,622,207) and Hauser, as applied to claims 68-74, 76 and 105 above, and further in view of Manning et al. (US 4,946,705).

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Iannocone in view of Moore, Wang and Hauser do not specifically teach that alumina particles are less than 180 μm in size.

Manning discloses "a simple, inexpensive, and versatile device for measuring gaseous substances and a method for manufacturing of such devices" (Abstract). The device comprises a reflectant baking on which are disposed microparticles coated at the surface with a reagent. "Suitable microparticulate adsorbents are particulate solids which have surface areas of at least 10 m²/g, preferably 100-1000 m²/g. These materials usually have a particle size of 100 microns or less, i.e., 0.1-100 microns. Materials which are available in this form include inorganic refractory oxidic supports, such as alumina, silica, glass, titania, clays and the like, or organic polymers which have been treated to achieve porosity, e.g., porous cellulose, polypropylene, and polystyrene-divinylbenzene particles. In order to be compatible with a detection reaction which generates color the support must be essentially colorless" (col. 7, lines 52-60). Therefore, ""microparticulate" support refers to particles which are reflectant, and finely divided so as to expose a large amount of surface area for coating with collection layers. The size of the particles is inversely proportional to the sensitivity of the device, since more surface area imparts more sensitivity to the test. Sizes of particles in the range capable of providing 10-1000 m²/g are suitable for uses in the invention, but this range is not taken to be limiting. Larger particle sizes are workable but decrease the sensitivity. Smaller particles are desirable from the standpoint of sensitivity but may be less convenient to handle. Any workable surface area/mass ratio is within the scope of the invention, and the suggested range is merely intended to provide guidance to practitioners thereof' (col. 4, lines 60-69).

It would have been obvious for anyone of ordinary skills in the art to use alumina particles of size less than 180 μ m, in particular of the size range of 0.1-100 microns, in Iannocone-Moore-Wang-Hauser's method of making the indicator for the same reason emphasized by Manning, i.e. to increase the surface area for coating and therefore increase the sensitivity of the indicator.

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Response to Arguments

12. Applicant's arguments filed 07/30/03 have been fully considered but they are not persuasive. The arguments are addressing the amended claims; however, even for amended claims the prior art makes the present invention obvious, as it discloses a method of making well-known acetaldehyde indicator based on MBTH method, with minor additional modifications of the structure of the indicator, which are also described in the prior art; applying the reagent-containing carrier to a support, or to an adhesive tape, which can be either double-sided thus providing an attachment area to any solid surface, or attaching one-sided adhesive tape to the solid surface using another piece of adhesive tape is well known to everyone of ordinary skill in the art and disclosed in the prior art cited above. Moore is not used as a primary reference in the present Office action, and therefore he does not teach away from modifying Iannocone's method; the modification of the method is related to providing a simple physical structure of the indicator as taught by Moore, and such structure can be employed for any indicator comprising dried reagent just for convenience of particular use.

Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP§706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event,

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however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Yelena G. Gakh, Ph.D. whose telephone number is (703) 306-5906. The examiner can normally be reached on 9:30 am - 6:00 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Jill A. Warden can be reached on (703) 308-4037. The fax phone number for the organization where this application or proceeding is assigned is (703) 872-9306.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-0661.

Yelena G. Gakh 9/15/03

Supervisory Patent Examiner Technology Center 1700